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RESULT 3
US-10-303-266-11
 Sequence 11, Application US/10303266
 Publication No. US20040101848A1
 GENERAL INFORMATION:
 APPLICANT: Donna T. Ward
 APPLICANT: Alexander H. Borchers
 APPLICANT: Kenneth W. Dobie
 TITLE OF INVENTION: MODULATION OF GLUCOSE TRANSPORTER-4 EXPRESSION
 FILE REFERENCE: RTS-0426
 CURRENT APPLICATION NUMBER: US/10/303,266
 CURRENT FILING DATE: 2002-11-23
 NUMBER OF SEQ ID NOS: 157
 SEQ ID NO 11
  LENGTH: 2128
  TYPE: DNA
  ORGANISM: H. sapiens
  FEATURE:
  NAME/KEY: CDS
  LOCATION: (146)...(1675)
US-10-303-266-11
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 Best Local Similarity 99.9%; Pred. No. 0;
 Matches 1529; Conservative
                     0; Mismatches
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         Db
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Qγ
         206 ACTGGGACCCTGGTCCTTGCTGTTCTCTGCGGTGCTTGGCTCCCTGCAGTTTGGGTAC 265
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      Qy
         Db
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Qν
         Db
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         926 AAGGATGAGAAGCGGAAGCTGGAGCGTGAGCGGCCACTGTCCCTGCTCCAGCTCCTGGGC 985
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          Db
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Dh
RESULT 4
US-10-067-449-9
; Sequence 9, Application US/10067449
Publication No. US20030166258A1
 GENERAL INFORMATION:
 APPLICANT: Muller, Gunter
  APPLICANT: Koller, Klaus-Peter
  APPLICANT: Boles, Eckhard
  APPLICANT: Wieczorke, Roman
  APPLICANT: Dlugai, Silke
 TITLE OF INVENTION: Saccharomyces cerevisiae Yeast Strain With Functional Expression of a TITLE OF INVENTION: GLUT Promoter
  FILE REFERENCE: DEAV2001/00002
  CURRENT APPLICATION NUMBER: US/10/067,449
  CURRENT FILING DATE: 2002-02-05
  PRIOR APPLICATION NUMBER: DE 101 06 718.6
  PRIOR FILING DATE: 2001-02-14
  NUMBER OF SEQ ID NOS: 18
  SOFTWARE: PatentIn version 3.0
 SEQ ID NO 9
  LENGTH: 7828
  TYPE: DNA
  ORGANISM: Homo sapiens
US-10-067-449-9
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                   99.9%; Score 1528.4; DB 6; Length 7828;
 Best Local Similarity
                   99.9%; Pred. No. 0;
 Matches 1529; Conservative
                        0; Mismatches
                                      1; Indels
        1 ATGCCGTCGGGCTTCCAACAGATAGGCTCCGAAGATGGGGAACCCCCTCAGCAGCGAGTG 60
Qy
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1: Biochim Biophys Acta, 1997 Feb 21;1324(1):111-9.

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Characterization of rat Glut4 glucose transporter expressed in the yeast Saccharomyces cerevisiae: comparison with Glut1 glucose transporter.

Kasahara T, Kasahara M.

Laboratory of Biophysics, School of Medicine, Teikyo University, Tokyo, Japan.

Rat Glut4 glucose transporter was expressed in the yeast Saccharomyces cerevisiae, but was retained in an intracellular membranous compartment and did not contribute to glucose uptake by intact cells. A crude membrane fraction was prepared and reconstituted in liposome with the use of the freeze-thaw/sonication method. D-glucose-specific, cytochalasin B inhibitable glucose transport activity was observed. Kinetic analysis of Dglucose transport was performed by an integrated rate equation approach. The K(m) under zero-trans influx condition was 12 +/- 1 mM (mean +/-S.E., n = 3) and that under equilibrium exchange condition was 22 +/- 3 mM (n = 4). D-glucose transport was inhibited by 2-deoxy-D-glucose or 3-Omethyl-D-glucose, but not by D-allose, D-fructose or L-glucose. Cytochalasin B, phloretin and phlorizin inhibited D-glucose transport, but neither p-chloromercuribenzoic acid (pCMB) (0-0.1 mM) nor pchloromercuribenzene sulfonic acid (pCMBS) (0-1.0 mM) inhibited this activity. High concentrations of HgCl2 were required to inhibit D-glucose transport (IC50, 370 microM). Comparing these properties to those of rat Glut1 we found two notable differences; (1) in Glut1, K(m) under zero-trans influx was significantly smaller than that under equilibrium exchange but in Glut4 less than two-fold difference was seen between these two K(m) values; and (2) Glut1 was inhibited with pCMB, pCMBS and low concentrations of HgCl2 (IC50, 3.5 microM), whereas Glut4 was almost insensitive to SH reagents. To examine the role of the exofacial cysteine, we replaced Met-455 of Glut4 (corresponding to Cys-429 of Glut1) with cysteine. The mutated Glut4 was inhibited by pCMB or pCMBS and the IC50 of HgCl2 decreased to 47 microM, whereas K(m), substrate specificity and the sensitivity to cytochalasin B were not significantly changed, indicating that the existence of exofacial cysteine contributed only to increase SH sensitivity in Glut4.